



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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<p>(21) International Application Number: PCT/EP99/01225</p> <p>(22) International Filing Date: 25 February 1999 (25.02.99)</p> <p>(30) Priority Data: MI98A000443 5 March 1998 (05.03.98) IT</p> <p>(71) Applicant (<i>for all designated States except US</i>): NICOX S.A. [FR/FR]; 45, avenue Kléber, F-75116 Paris (FR).</p> <p>(72) Inventor; and</p> <p>(75) Inventor/Applicant (<i>for US only</i>): DEL SOLDATO, Piero [IT/IT]; Via Toti, 22, I-20052 Monza (IT).</p> <p>(74) Agents: SAMA, Daniele et al.; Sama Patents, Via G.B. Morgagni, 2, I-20129 Milano (IT).</p>		<p>(81) Designated States: AL, AU, BB, BG, BR, CA, CN, CZ, EE, GE, HU, IL, IS, JP, KP, KR, LK, LR, LT, LV, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, TR, TT, UA, US, UZ, VN, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).</p> <p>Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i></p> <p>(88) Date of publication of the international search report: 25 November 1999 (25.11.99)</p>	

(54) Title: PHARMACEUTICAL COMPOSITIONS FOR ULCER

(57) Abstract

Pharmaceutical compositions for the therapy and the prevention of ulcer and dyspepsia relapses comprising as essential components the following components a) and b): component a): one or more components selected from the conventional anti-ulcer products; component b): organic compounds containing the -ONO₂ function, or inorganic compounds containing the -NO group, characterized in that they are NO nitric oxide donors.

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INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 99/01225

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 6 A61K33/00 A61K31/415 // (A61K33/00, 31:415)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 97 31654 A (NICOX SA ;DEL SOLDATO PIERO (IT)) 4 September 1997 see abstract see page 1 - page 5 see table 5 see claims 1-12	1-20
Y	---	1-20 -/-



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

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Date of the actual completion of the international search

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INTERNATIONAL SEARCH REPORT

International Application No PCT/EP 99/01225

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	HISANAGA, Y. ET AL.: "Implication of nitric oxide synthase activity in the genesis of water immerstion-stress induced gastric lesions in rats: the protective effects of FK-506" ALIMENT. PHARMACOL. THER., vol. 10, 1996, pages 933-940, XP002108980 see abstract see tables 1,2 see figures 1,2 see Discussion, especially the last two paragraphs ---	1-20
P,X	KIUCHI Y., ET AL.: "Effect of pibutidine hydrochloride on nitric oxide production in rat gastric mucosa" RES. COMM. MOL. PATH. PHARMACOL., vol. 100, no. 3, June 1998, pages 273-281, XP002108981 see abstract see Results see Discussion -----	1-20

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 99/01225

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO 9731654	A 04-09-1997	IT	MI960352 A	26-08-1997
		AU	706591 B	17-06-1999
		AU	2092497 A	16-09-1997
		CA	2247848 A	04-09-1997
		EP	0904110 A	31-03-1999

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(71) Applicant (<i>for all designated States except US</i>): NICOX S.A. [FR/FR]; 45, avenue Kléber, F-75116 Paris (FR). (72) Inventor; and (75) Inventor/Applicant (<i>for US only</i>): DEL SOLDATO, Piero [IT/IT]; Via Toti, 22, I-20052 Monza (IT). (74) Agents: SAMA, Daniele et al.; Sama Patents, Via G.B. Morgagni, 2, I-20129 Milano (IT).			
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PHARMACEUTICAL COMPOSITIONS FOR ULCER

The present invention relates to compositions to be used in the therapy and in the prevention of the ulcer relapses and, in general of the dyspepsias. More particularly it relates to compositions having an improved gastroprotective activity combined with an high acid secretion inhibition activity.

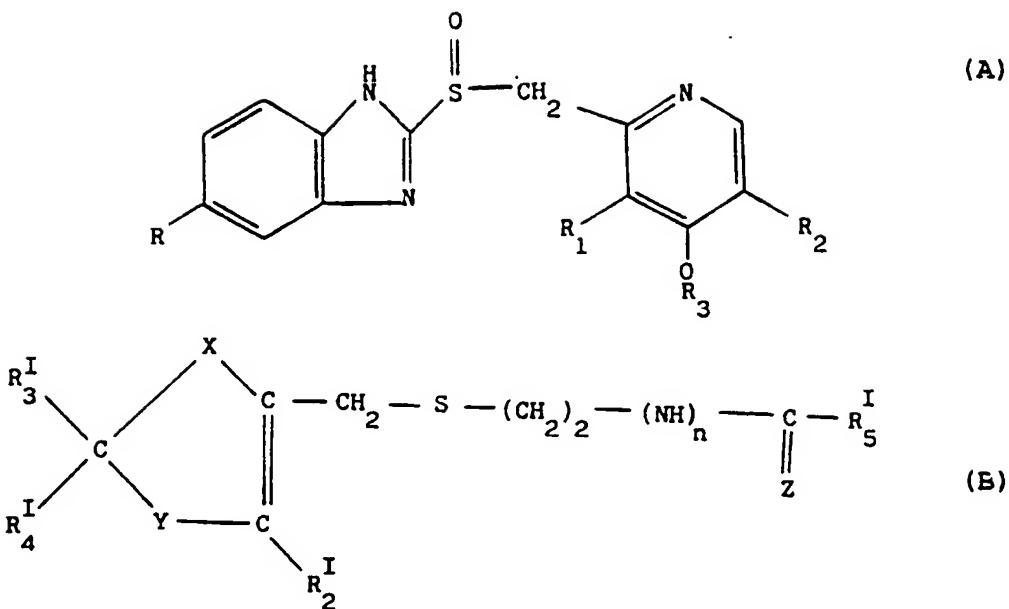
The products known in the art and those commercialized and used in the ulcer therapy are compounds which perform an anti-secretory activity (acid secretion inhibition). See for instance "New Guide to Medicine & Drugs" Brit. Medical Assoc. Editor, 1997, pages 108-109. Known products having higher therapeutic efficacy show an high anti-secretory activity and are used, both in the acute and in the long-term (6 months and more) therapies. The drawback of these products is that they have a poor gastroprotective activity, when present. From a practical point of view this means that the gastric protection is not optimal and causes inconveniences especially in the long-term therapy. In this case the presence of frequent relapses due to the enfeeblement of gastric mucosa is noticed.

To overcome these inconveniences it is known in the art to add to said medicaments other anti-ulcer medicaments having a gastroprotective action: prostaglandins, bismuth salts (e.g. bismuth citrate) and antibiotics. In such way the removal of ulcerous pathology is achieved. However these combinations are not satisfactory as to their tolerability in general. For example it is well known that prostglandins produce side effects (diarrhoea) towards the intestinal tract; bismuth salts frequently produce nausea and heartburn. Antibiotics produce undesired gastrointestinal effects.

The need was felt to have available compositions active in the ulcer and gastric dyspepsia treatment having improved therapeutic characteristic and tolerability, general and local, in particular having an improved gastroprotective activity combined with an high anti-secretion activity.

The Applicant has unexpectedly and surprisingly found pharmaceutical anti-ulcer compositions having an improved therapeutic activity and tolerability.

It is an object of the present invention pharmaceutical compositions comprising as essential components the following: component a): one or more components selected from the following classes:



wherein in the (A) class compounds

$R = H, OCH_3, OCHF_2;$

$R_1 = CH_3, OCH_3;$

$R_2 = H, CH_3;$

$R_3 = CH_3, CH_2-CF_3, (CH_2)_3-OCH_3;$

wherein in the (B) class compounds:

R_3^I, R_4^I equal to or different from each other, are respectively free valence, hydrogen, $-N=C(NH_2)_2, -CH_2-N(CH_3)_2;$

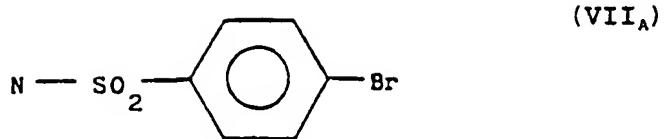
$Y = S, N-R_6^I, CR_7^I R_8^I;$

$X = O, S, N-R_1^I;$

$R_2^I = H, CH_3;$

$n = 0, 1;$

$Z = N-CN, N-SO_2NH_2, CH-NO_2$ or



$R^1_5 = H, -NH-CH_3, NH_2;$

$R^1_6, R^1_7, R^1_8, R^1_9$, equal to or different from each other, are hydrogen, free valence;

component b): one or more organic compounds containing the -ONO₂ function, or one or more inorganic compounds containing the -NO group, or mixtures thereof, said compounds being characterized in that they are NO nitric oxide donors, i.e. when they are brought into contact in vitro with cells of the basal endothelium, platelets, etc., and after incubation of 5 minutes at a temperature of 37°C, are capable to release NO and to activate the cGMP synthesis (Guanosine cyclic 3',5'-(hydrogen phosphate)), as shown by the used specific tests which will be described in detail further on.

The present invention compositions are obtainable by directly mixing the components a) and b).

The molar ratio between the components a) and b) can vary within wide limits, from 1 : 0.1 to 1 : 5, preferably from 1 : 0.5 to 1 : 2.

As regards the component a), the preferred (A) com-

pounds are the following:

when R = OCH₃, R₁ = CH₃, R₂ = CH₃, R₃ = CH₃, Omeprazole residue;

as in Omeprazole, but with R = OCHF₂, R₁ = OCH₃, R₂ = H, Pantoprazole residue;

as in Omeprazole, but with R = H, R₂ = H, R₃ = (CH₂)₂-OCH₃, Rabeprazole residue;

as in Rabeprazole, but with R₁ = CH₂-CF₃, Lansoprazole residue;

in component a) the preferred compounds of formula (B) are the following:

when in formula (B) X = N-R^I₁, with R^I₁ free valence, Y = N-R^I₆ with R^I₆ = H, R^I₃ = H, R^I₄ is free valence and forms with R^I₁ a double bond, R^I₂ = CH₃, n = 1, R^I₅ = -NH-CH₃, Z = N-CN, Cimetidine residue;

when X = N-R^I₁ with R^I₁ free valence, Y = S, R^I₃ = -N=C(NH₂)₂, R^I₄ is free valence and forms with R^I₁ a double bond, R^I₂ = H, n = 1 R^I₅ = H, Z = (VII_A), Ebrotidine residue;

as in Ebrotidine but with n = 0, R^I₅ = NH₂ and Z = N-SO₂NH₂, Famotidine residue;

as in Ebrotidine but with R^I₁ = -CH₂-N(CH₃)₂, R^I₅ = -NH-CH₃ and Z = CH-NO₂, Nizatidine residue;

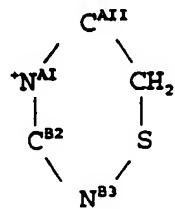
as in Nizatidine, but with X = oxygen, Y = CR^I₁R^I₆ with R^I₁ hydrogen and R^I₆ free valence, R^I₄ is a free valence and forms

with Rⁱ, a double bond, Ranitidine residue.

In the preparation of the invention compositions also the isomers of one or more compounds of each component can be used.

The components belonging to (A) and (B) classes are prepared according to methods described in "The Merck Index 12^a Ed." (1996), herein incorporated by reference.

In the formula (A) compounds of component a) also the compounds having the following intramolecular ring are comprised, obtainable by treating the precursors in an acid aqueous medium (ref. " A Textbook of Drug Design and Development", Harwood academic publisher, 1991, pag. 140):



wherein N^{A1} and C^{A11} are, respectively, the nitrogen and the carbon atom in 1 and 2 position of the pyridine ring of formula (A) and C^{B2} and N^{B3} the carbon and nitrogen atom, respectively, in 2 and 3 position of the imydazole ring (1 position of the imydazole ring is that of the proton nitrogen).

Instead of the formula (A) and (B) compounds of component a) the corresponding salts with organic or inorganic salts, obtainable by reactions known to the skilled in the

art, can be used. Examples of usable inorganic ions are the following: hydrochlorides, nitrates, sulphates, phosphates, etc.; among the organic ones maleates, oxalates, acetates, succinates, etc. can be mentioned. Compositions containing nitrates are preferred.

The salts with the preferred compounds of the component a) are those corresponding to the preferred compounds of formula (A) or (B).

In the compositions according to the present invention salts of the compound of the above formulas contain at least one anion mole/compound mole. Preferably the ratio between the anion and compound moles is unitary. Salts having a higher molar ratio are obtained when in the molecule other basic enough aminic groups to be salified are present.

For example the salts of the class (A) and (B) compound with nitric acid are prepared according to the following methods.

When the substance to be salified is available as free base or as a corresponding salt soluble in an organic solvent, which preferably does not contain hydroxyl groups, for example acetonitrile, ethyl acetate, tetrahydrofuran, etc., the salt is prepared by dissolving the substance in the solvent at a concentration preferably equal or higher than 10% w/v, by adding the amount of concentrated nitric acid cor-

responding to the moles of salifiable aminic groups present in the compound. The nitric acid is preferably diluted in the same solvent. Preferably during and after the addition the mixture is cooled to temperatures in the range 20°C-0°C. The product is generally recovered by filtration and washed with the solvent.

When on the contrary the substance is not very soluble, or it is available as a not very soluble salt in the above mentioned solvents, the corresponding mixtures with hydroxylated solvents may be used. Examples of such solvents are methyl alcohol, ethyl alcohol and water. Precipitation can be quickened by diluting then the so obtained mixture, after the addition of nitric acid, with an apolar solvent.

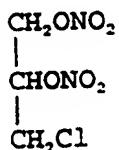
When the starting product is salified with hydrochloric acid, it is possible to prepare the nitric acid salt directly adding silver nitrate to the solution. After filtering silver chloride, the solution is concentrated and treated as above indicated to recover the nitrate salt.

When the starting product is a salt, it is also possible to liberate the corresponding base by a treatment with a sodium or potassium bicarbonate or carbonate saturated solution, or with a sodium or potassium hydroxide diluted solution. The base is then extracted by a suitable organic solvent (for example halogenated solvents, esters, ethers), which is then dried. The organic solution is evaporated and

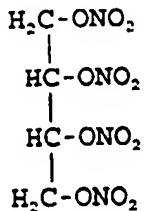
then one proceeds according to the preceding preparation methods, by dissolving the base in acetonitrile or in the other above mentioned solvents.

For the component b), the organic compounds containing the -ONO₂ groups which can be used are, as an example, the following, which are reported in The Merck Index 11th Ed. 1989 and prepared by the known methods, for instance those mentioned in the same reference, herein incorporated by reference:

chloronitrate (3-chloro-1,2-propanediol dinitrate) (Merck No. 2390) having the formula C₃H₅ClN₂O₆ and structural formula

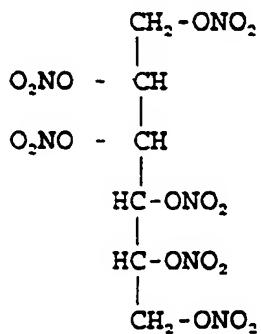


erythrityltetranitrate (1,2,3,4 butantetroltetranitrate) (Merck No. 3622) having the formula C₄H₆N₄O₁₂ and structural formula



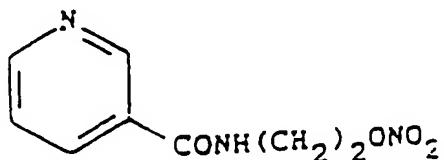
mannitol hexanitrate (Merck No. 5630) having the formula

C₈H₉N₆O₁₈ and structural formula

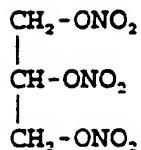


nicorandyl (N-[2-(nitroxy)ethyl]-3-pyridine-carboxamide)

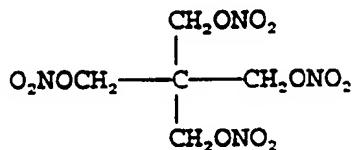
(Merck No. 6431) having the formula C₈H₉N₆O₄ and structural formula



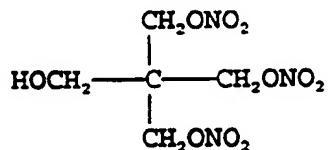
nitroglycerin (1,2,3 propanetriol trinitrate) (Merck No. 6528) having the formula C₃H₅N₁O₉, and structural formula



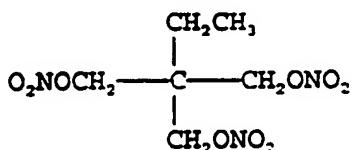
pentaerythritoltetranitrate (2,2-bis [(nitroxy)-methyl]-1,3-propanedioldinitrate) (Merck No. 7066) having the formula C₅H₆N₄O₁₂ and structural formula



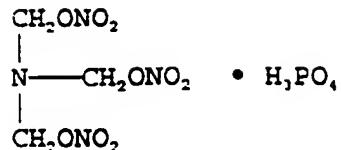
pentrinitrol (2,2-bis[(nitroxy)methyl]-1,3-propanediolmononitrate) (Merck No. 7094) having the formula C₅H₉N₃O₁₀ and structural formula



propetrylnitrate (2-ethyl-2-[(nitroxy)methyl]-1,3-propanedioldinitrate) (Merck No. 7821) having the formula C₆H₁₁N₃O₉ and structural formula



trolnitratephosphate (2,2',2"-nitryltrisethanoltrinitrate phosphate) (1:2 salt) (Merck No. 9682) having the formula C₆H₁₈N₄O₁₇P₂ and structural formula



Other compounds containing the -ONO₂ group which can be used are prostglandin nitroderivatives described in the PCT/EP Application 98/03645 in the Applicant's name. This patent application is herein incorporated by reference both for compounds and preparation processes.

Among the inorganic compounds containing the -NO group can be mentioned:

- nitrate salts of alkaline, earth-alkaline or III group metals, for example with aluminum, sodium and potassium nitrates are preferred;
- nitroprussiate salts, such as for example: sodiumnitroprussiate (pentakis (cyano-C)nitrosylferrate(2-)disodium) (Merck No. 8600), having the formula Na₂[Fe(CN)₅NO].

The invention components b) containing the -ONO₂ groups or the -NO group which produce the invention results, as already said, must meet the test in vitro described in detail hereinunder.

Specifically the test relates to the nitric oxide generation from the NO-donors described in the present invention, among which, for instance nitroglycerin, nicorandyl, nitroprussiate, etc., when they are put in the presence of

endothelial cells (method a) or platelets (method b):

a) Endothelial cells

Cells of the human umbilical vein, spread on plate, with a 10^3 cells/plate density have been incubated with scalar concentrations of NO-donor (1-100 $\mu\text{g}/\text{ml}$) for 5 minutes. The incubation medium (physiologic solvent, for example Tyrode) has been then analyzed to determine the capability to generate NO, by:

- 1) the nitric oxide disclosure by chemiluminiscence;
- 2) cGMP determination (cyclic GMP No. 2715 of the above mentioned Merck).

As regards the chemiluminiscence analysis, an amount equivalent to 100 μl was injected in the reaction chamber of a chemiluminiscence analyzer containing glacial acetic acid and potassium iodide. Nitrites/nitrates present in the medium under these conditions are converted into NO which is then determined after its reaction with ozone, with consequent light generation. As it usually occurs in the devices measuring the chemiluminiscence, the produced luminiscence is directly proportional to the generated NO levels and can be measured by the suitable photomultiplier unit of a chemiluminiscence analyzer.

The photomultiplier converts the incident light into electric voltage, which is then quantitatively re-

corded. On the basis of a calibration curve, prepared with scalar nitrite concentrations, it was possible to quantitatively determine the generated NO concentration. For example, from the incubation of 100 μM of nicorandyl, an amount equal to about 10 μM of NO was generated.

As regards the cGMP determination, a portion of the incubation medium (equal to 100 μl) was centrifuged at 1000 revolutions per 20 seconds. The supernatant was discharged and the sediment treated with iced phosphate buffer (pH 7,4). The cGMP level produced were tested, by specific immuno-enzymatic reactants. From such experiments it resulted that, under these experimental conditions, the incubation with one of the various tested NO-donors caused a significant cGMP increase with respect to the values obtained in absence of a NO donor. For example, after incubation with 100 μM of sodium nitroprussiate, an increase of about 20 times the value obtained with the incubation of only the vehicle without NO donor was recorded.

b) Platelets

Washed human platelets, prepared analogously with what described by Radomski et al, (Br.J. Pharmacol. 92, 639-1987), were used. 0.4 ml aliquots were incubated with scalar NO-donor concentrations (1-100 $\mu\text{g/ml}$) for 5 mi-

nutes. The incubation medium (e.g. Tyrode) was then analyzed to determine the capability to generate NO, by determination of nitric oxide by chemiluminiscence and cGMP, with the modalities described in the previous paragraph, for the analyses carried out on the endothelial cells. As to the determination by chemiluminiscence, also in this case, on the basis of a calibration curve, prepared with scalar nitrite concentrations, it was possible to quantitatively determine the generated NO concentration. For example, after incubation of 100 μM of nicorandyl, an amount equal to 35 μM of NO was generated.

As regards the cGMP determination, also in these experimental conditions it resulted that the incubation with one of the various NO-donors tested caused a cGMP significant increase with respect to the values obtained in absence of a NO-donor. For example, after incubation with 100 μM of sodium nitroprussiate, an increase of about 30 times the value obtained with the incubation of only the vehicle without NO-donor was recorded.

In conclusion, from said tests it results that all the NO-donors according to the present invention, after incubation with endothelial cells or platelets for 5 minutes, are capable to generate NO, and to activate the cGMP synthesis

in a concentration-depending way, as determined by the used specific tests.

It has now surprisingly been found that the present invention compositions allow to improve, compared with the known above mentioned combinations, the total pharmaco-toxicological behaviour of the compounds forming the component a), increasing the therapeutic efficacy and their general and local tolerability in the ulcer and gastric dyspepsia treatment, with improved gastroprotective activity.

The present invention compositions are formulated in the corresponding pharmaceutical compositions according to well known techniques in the field together with the conventional excipients; see for example the volume "Remington's Pharmaceutical Sciences 15a Ed."

The administering of the present invention compositions can be carried out by oral, parenteral or transdermic way. The component b) can generally be administered contemporaneously, subsequently or previously to the component a).

The (A) and (B) compound doses of component a) are the conventional ones for these compounds.

It is a further object of the invention compositions obtainable by combining the pharmaceutical compositions according to the present invention with conventional gastroprotective agents. As examples, prostaglandins, bismuth salts, active antibiotics towards pathogenic microorganisms

in the gastrointestinal mucosa can be mentioned.

It has surprisingly been found that the gastroprotective activity of the invention compositions is very high. This makes it possible to avoid the undesired effects of known gastroprotectives when they are used in combination with compounds or formulations of the invention compounds. It has indeed been found that the amount of the known gastroprotective agents, in the combination of the invention, is lower compared with those known and does not cause undesirable side effects. The skilled in the field is able to easily determine the maximum amount of conventional gastroprotective agents to be combined with the invention pharmaceutical compositions, since this corresponds to the absence of typical side effects of the known gastroprotective agents. In any case the amount of conventional gastroprotective agents to be used in the combination is lower than that used in combinations described in the art.

The following examples have the purpose to illustrate the invention and must not be considered as limitative of the same.

EXAMPLE 1

Pharmaceutical composition.

Mixture formed by:

Component a) cimetidine 50 g.

component b) KNO₃ 20 g.

Amounts of this mixture corresponding to 70 mg/kg, or multiples, to be supplied to animals, are then weighed.

Pharmacological Tests

EXAMPLE 2

Acute toxicity

To a group of 10 rats weighing 20 g each a single dose equal to 280 mg/kg of the Example 1 composition was supplied by oral way in a carboxymethylcellulose aqueous suspension 2% w/v.

The animals are kept under obsevation for 14 days. In no one of the group animals the toxic syntom presence was noticed.

EXAMPLE 3

Anti-ulcer Activity

The anti-ulcer activity is evaluated according to the experimental model described in the paper of A. Robert et Al. "Cytoprotection by prostaglandins in rats. Prevention of gastric necrosis produced by alcohol, HCl, NaOH, hypertonic NaCl and thermal injury" Gastroenterology 77, 433-43 1979.

To 3 groups of 10 rats each, kept with empty stomach since the previous night, 15 minutes before the supply of absolute ethanol (1 ml), by oral way are supplied:

- 5 ml/Kg of carboxymethylcellulose aqueous suspension 2%.
- 50 mg/Kg of cimetidine in 5 ml/Kg of carboxymethylcel-

lulose aqueous suspension 2%.

- 70 mg/Kg of the Example 1 composition in 5 ml/Kg of carboxymethylcellulose aqueous suspension 2%.

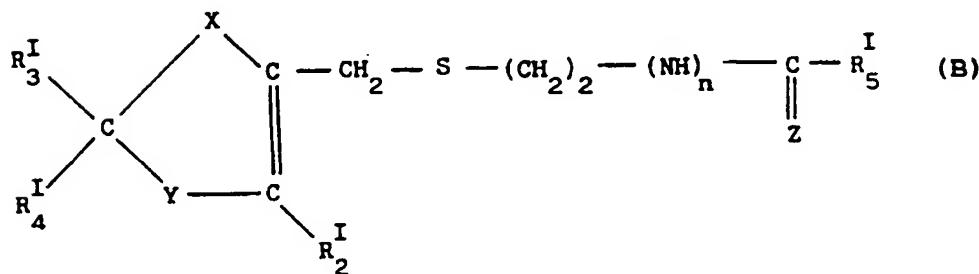
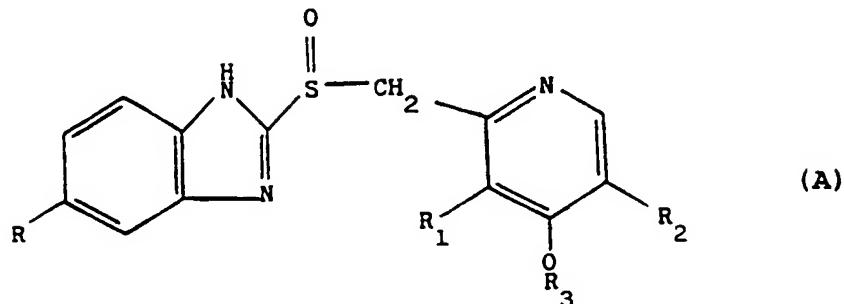
One hour later the animals are sacrificed and the gastric lesion incidence is evaluated. The results are reported in Table 1 and they show that the combination cimetidine and potassium nitrate shows an improved gastroprotective activity with respect to cimetidine.

TABLE I

Treatment	Gastric Damage (%)
Vehicle	100
Cimetidine 50mg/Kg	100
Composition Ex. 1	
70 mg/Kg	60

CLAIMS

1. Pharmaceutical compositions comprising as components
the following:
component a): one or more components selected from the
following classes:



wherein in the (A) formula

R = H, OCH₃, OCHF₂;

R₁ = CH₃, OCH₃;

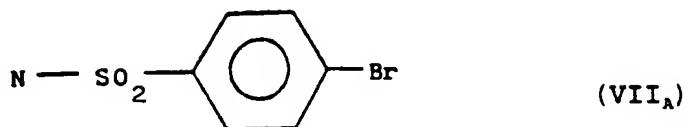
R₂ = H, CH₃;

R₃ = CH₃, CH₂-CF₃, (CH₂)₃-OCH₃;

in the (B) class compounds:

R^I₃, R^I₄ equal to or different from each other, are re-

spectively free valence, hydrogen, -N=C(NH₂)₂,
 -CH₂-N(CH₃)₂; Y = S, N-R^I₆, CR^I₂R^I₅;
 X = O, S, N-R^I₁;
 R^I₂ = H, CH₃;
 n = 0, 1;
 Z = N-CN, N-SO₂NH₂, CH-NO₂ or



R^I₅ = H, -NH-CH₃, NH₂;

R^I₆, R^I₂, R^I₃, R^I₄, equal to or different from each other,

are hydrogen, free valence;

component b): one or more organic compounds containing the -ONO₂ group, or one or more inorganic compounds containing the -NO group, or mixtures thereof, said compounds being characterized in that they are NO nitric oxide donors, i.e. when they are brought into contact in vitro with cells of the basal endothelium, platelets, and after incubation of 5 minutes at a temperature of 37°C, are capable to release NO and to activate

the cGMP synthesis (Guanosine cyclic 3',5'-(hydrogen phosphate)).

2. Pharmaceutical compositions according to claim 1, wherein the molar ratio between the components a) and b) varies from 1 : 0.1 to 1 : 5, preferably from 1 : 0.5 to 1 : 2.
3. Pharmaceutical compositions according to claims 1-2 wherein in component a) the compounds are the following:

- formula (A) :

R = OCH₃, R₁ = CH₃, R₂ = CH₃, R₃ = CH₃, Omeprazole residue;

as in Omeprazole, but with R = OCHF₂, R₁ = OCH₃, R₂ = H, Pantoprazole residue;

as in Omeprazole, but with R = H, R₂ = H, R₃ = (CH₂)₃-OCH₃, Rabeprazole residue;

as in Rabeprazole, but with R₃ = CH₂-CF₃, Lansoprazole residue;

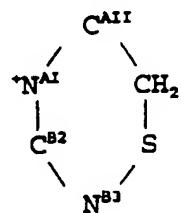
- formula (B) :

X = N-R^I₁ with R^I₁ free valence, Y = N-R^I₆ with R^I₆ = H, R^I₄ = H, R^I₄ is free valence and forms with R^I₁ a double bond, R^I₂ = CH₃, n = 1, R^I₅ = -NH-CH₃, Z = N-CN, Cimetidine residue;

when in the formula (B) X = N-R^I₁ with R^I₁ free

valence, Y = S, R^I₃ = -N=C(NH₂)₂, R^I₄ is free valence and forms with R^I₁ a double bond, R^I₂ = H, n = 1 R^I₅ = H, Z = (VII_A), Ebrotidine residue; as in Ebrotidine but with n = 0, R^I₅ = NH₂ and Z = N-SO₂NH₂, Famotidine residue; as in Ebrotidine but with R^I₃ = -CH₂-N(CH₃)₂, R^I₅ = -NH-CH₃, and Z = CH-NO₂, Nizatidine residue; as in Nizatidine, but with X = oxygen, Y = CR^I₂R^I₃ with R^I₁ hydrogen and R^I₂ free valence, R^I₄ is a free valence and forms with R^I₆ a double bond, Ranitidine residue.

4. Pharmaceutical compositions according to claims 1-3, wherein the isomers of one or more compounds of each component are used.
5. Pharmaceutical compositions according to claims 1-4, wherein in the formula (A) compounds of component a) also the compounds having the following intramolecular ring are comprised, obtainable by treating the precursors in acid aqueous medium:



wherein N^{A1} and C^{A1} are, respectively, nitrogen and carbon atoms in 1 and 2 position of the pyridine ring and C^{B2} and N^{B3} the carbon and nitrogen atoms, respectively, in position 2 and 3 of the imydazol ring.

6. Pharmaceutical compositions according to claims 1-5, wherein instead of the compounds of formula (A) and (B) of component a) the corresponding organic or inorganic acid salts can be used.
7. Pharmaceutical compositions according to claim 6 containing nitrate salts.
8. Pharmaceutical compositions according to claim 6 containing the salts of formula (A) or (B) compounds of claim 3.
9. Pharmaceutical compositions according to claims 6-8, wherein the salts contain at least one anion mole/compound mole.
10. Pharmaceutical compositions according to claims 1-9, wherein the organic compounds containing the -ONO₂ groups usable in component b) are selected from the following ones:
clonitrate, erytrityltetranitrate, mannitol hexanitrate, nicorandyl, nitroglycerin, pentaerythritoltetrinitrate, pentrinitrol, propatylnitrate trolnitratephosphate, prostaglandin nitroderivatives.
11. Pharmaceutical compositions according to claims 1-9 wh -

rein the inorganic compounds containing the -NO group of component b) are selected from the following: nitrate salts of alkaline, earth-alkaline or III group metals, and nitroprussiate salts.

12. Pharmaceutical compositions according to claim 11 wherein the nitrate salts are sodium and potassium nitrate.
13. Pharmaceutical compositions according to the previous claims formulated for the oral, parenteral, transdermic administering.
14. Pharmaceutical compositions according to claim 13 wherein the component b) is administered contemporaneously, subsequently or previously to component a).
15. Pharmaceutical compositions according to claims 1-14 for use as a medicament.
16. Use of the pharmaceutical compositions according to claims 1-14, to prepare medicaments for the therapy and prevention of the ulcer and dyspepsia relapses.
17. Use according to claims 15-16 wherein the (A) and (B) compound doses of component a) are the conventional ones used for these compounds.
18. Pharmaceutical compositions obtainable by combining the compositions according to claims 1-14 with conventional gastroprotective agents.
19. Compositions according to claim 18 wherein the conven-

tional gastroprotective agents are selected from pro-
staglandins, bismuth salts and antibiotics.

20. Use of pharmaceutical compositions according to claims
18-19 to prepare medicaments for the therapy and pre-
vention of the ulcer and dyspepsia relapses.